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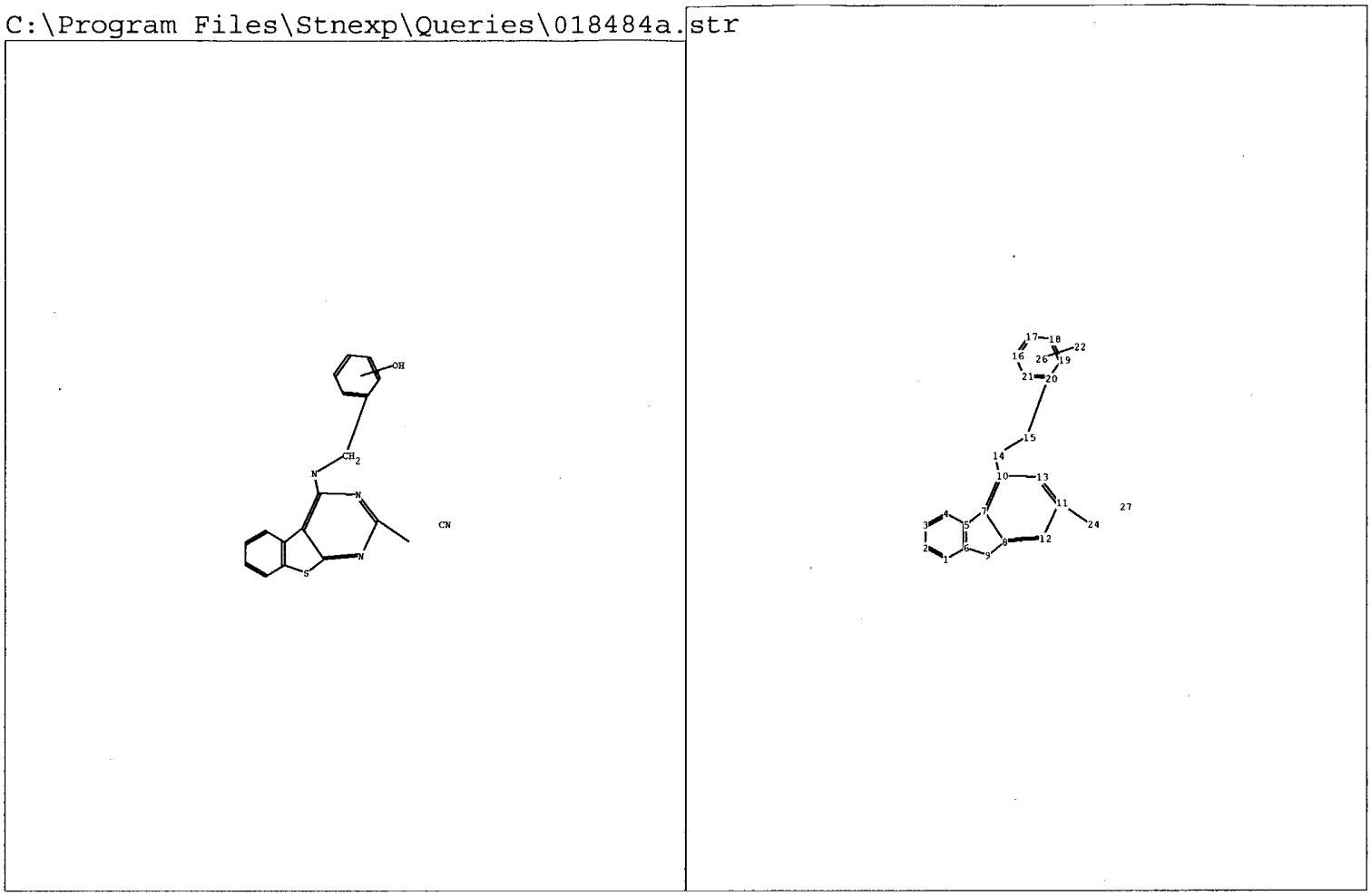

chain nodes :
14 15 22 27
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 16 17 18 19 20 21
ring/chain nodes :
24
chain bonds :
10-14 11-24 14-15 15-20
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 7-10 8-9 8-12 10-13
11-12 11-13 16-21 16-17 17-18 18-19 19-20 20-21
exact/norm bonds :
10-14
exact bonds :
5-7 6-9 8-9 11-24 14-15 15-20
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-10 8-12 10-13 11-12 11-13 16-21
16-17 17-18 18-19 19-20 20-21
isolated ring systems :
containing 1 : 16 :

G1:O,N

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom
10:Atom 11:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 16:Atom 17:Atom
18:Atom

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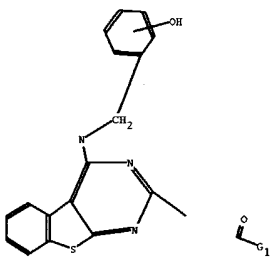

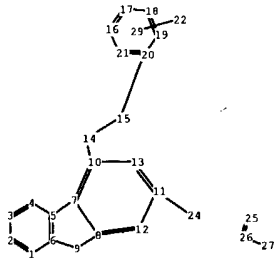

chain nodes :
14 15 22 25 26 27
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 16 17 18 19 20 21
ring/chain nodes :
24
chain bonds :
10-14 11-24 14-15 15-20 25-26 26-27
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 7-10 8-9 8-12 10-13
11-12 11-13 16-21 16-17 17-18 18-19 19-20 20-21
exact/norm bonds :
10-14 25-26 26-27
exact bonds :
5-7 6-9 8-9 11-24 14-15 15-20
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-10 8-12 10-13 11-12 11-13 16-21
16-17 17-18 18-19 19-20 20-21
isolated ring systems :
containing 1 : 16 :

G1:O,N

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom
10:Atom 11:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 16:Atom 17:Atom
18:Atom

Property values tagged with IC are from the ZIC/VINITI data file
provided by InfoChem.

STRUCTURE FILE UPDATES: 22 APR 2004 HIGHEST RN 676515-68-7
DICTIONARY FILE UPDATES: 22 APR 2004 HIGHEST RN 676515-68-7

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more
information enter HELP PROP at an arrow prompt in the file or refer
to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

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L1 STRUCTURE UPLOADED

=>
Uploading C:\Program Files\Stnexp\Queries\018484.str

L2 STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> d l2
L2 HAS NO ANSWERS
L2 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

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FULL SEARCH INITIATED 12:28:18 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 3 TO ITERATE

100.0% PROCESSED 3 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

L3 0 SEA SSS FUL L1

=> end

END`````````````````` IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).

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FULL SEARCH INITIATED 12:28:35 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 86 TO ITERATE

100.0% PROCESSED 86 ITERATIONS 6 ANSWERS
SEARCH TIME: 00.00.01

L4 6 SEA SSS FUL L2

=> file caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	310.84	311.05

FILE 'CAPLUS' ENTERED AT 12:28:44 ON 24 APR 2004
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

FILE COVERS 1907 - 24 Apr 2004 VOL 140 ISS 18
FILE LAST UPDATED: 23 Apr 2004 (20040423/ED)

This file contains CAS Registry Numbers for easy and accurate
substance identification.

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L5 5 L4

=> d l5 1-5 ibib abs hitstr

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:695578 CAPLUS
DOCUMENT NUMBER: 137:206584
TITLE: Oral drug delivery systems containing benzothieno-pyrimidines
PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany
SOURCE: Ger. Offen., 8 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10107261	A1	20020912	DE 2001-10107261	20010216
WO 2002072100	A2	20020919	WO 2002-EP609	20020123
WO 2002072100	A3	20031106		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EE 200300378	A	20031015	EE 2003-378	20020123
EP 1385521	A2	20040204	EP 2002-701263	20020123

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

BR 2002007271	A	20040323	BR 2002-7271	20020123

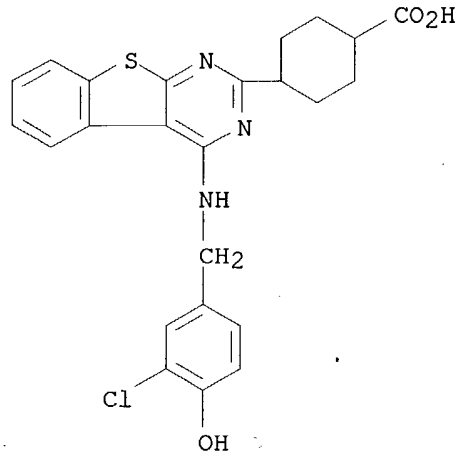
PRIORITY APPLN. INFO.: DE 2001-10107261 A 20010216
WO 2002-EP609 W 20020123

AB The invention concerns a pharmaceutical composition with high bioavailability for the oral administration of 4-[4-(3-Chloro-4-methoxy-benzylamino)-benzo[4,5]thieno[2,3-d]pyrimidin-2-yl] cyclohexane carboxylic acid and/or 4-[4-(3-Chloro-4-hydroxy-benzylamino)-benzo[4,5]thieno[2,3-d]pyrimidin-2-yl]-cyclohexane carboxylic acid or their pharmaceutically acceptable salts that contain a surfactant with HLB value 14-16.7 and another surfactant with HLB value 3-5. Thus the content of a soft gelatin capsule was prepared from (mg): 4-[4-(3-Chloro-4-methoxy-benzylamino)-benzo[4,5]thieno[2,3-d]pyrimidin-2-yl] cyclohexane carboxylic acid ethanolamine salt 50; polyethylene-hydrated castor oil 180; medium-chain partial glycerides 135; PEG 400 90; glycerol (85%) 45.

IT **313641-90-6**, Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]-
RL: PEP (Physical, engineering or chemical process); PYP (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
(pharmaceutical composition containing benzothieno-pyrimidines)

RN 313641-90-6 CAPLUS

CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)

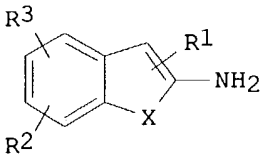


REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:762987 CAPLUS
DOCUMENT NUMBER: 135:303761
TITLE: Preparation of benzo-annelated heterocycles by aromatization of tetrahydrobenzo-annelated heterocycles with a hydrogenation catalyst in the presence of a hydrogen acceptor
INVENTOR(S): Wartenberg, Frank-Hardi; Koppe, Thomas; Wetzel, Walter; Wydra, Markus; Benz, Achim
PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany
SOURCE: PCT Int. Appl., 26 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001077099	A1	20011018	WO 2001-EP2672	20010309
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
DE 10017947	A1	20011025	DE 2000-10017947	20000411
EP 1280792	A1	20030205	EP 2001-921324	20010309
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
BR 2001009937	A	20030527	BR 2001-9937	20010309
JP 2003530392	T2	20031014	JP 2001-575572	20010309
US 2003078422	A1	20030424	US 2002-257139	20021009
US 6723848	B2	20040420		
NO 2002004887	A	20021010	NO 2002-4887	20021010

PRIORITY APPLN. INFO.: DE 2000-10017947 A 20000411
WO 2001-EP2672 W 20010309
OTHER SOURCE(S): CASREACT 135:303761; MARPAT 135:303761
GI



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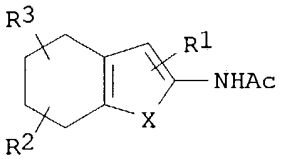


II

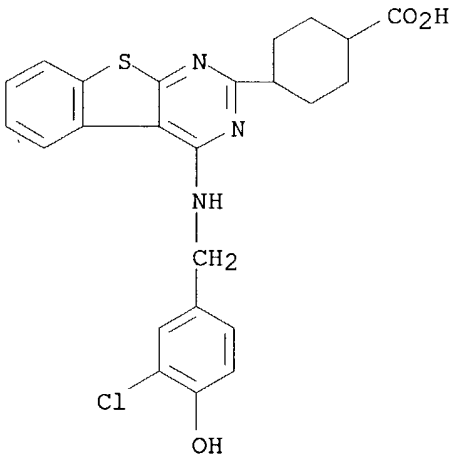
AB Title compds. [I; X = S, O, NH; R1 = cyano, NO2, acyl, CO2H, CO2A, CO2NR4R5, (substituted) PhCO, CO2Ph; R2, R3 = H, alkyl, NO2, cyano, OH, alkyloxy, acyl; R4, R5 = H, alkyl, acyl, (substituted) Ph; R4R5 = CH2(CH2)nCH2; A = alkyl; n = 2-4] are prepared by reaction of II (R1-R3 as

above, Ac = acyl) with a noble metal catalyst in the presence of a H acceptor followed by deacylation of the acylated amino group with an amine. Thus, 2-acetylamino-4,5,6,7-tetrahydrobenzothiophene-3-carboxylic acid Et ester in mesitylene was heated at 170° with Pd/C under stirring followed by addition of Et cinnamate over 30 min. stirring for 21 h, and addition at 100° of pyrrolidine to give after 25 h stirring 64% 2-aminobenzo[b]thiophene-3-carboxylic acid Et ester. The latter is an intermediate for 4-[4-(3-chloro-4-methoxybenzylamino)benzo[4,5]thieno[2,3-d]pyrimidin-2-yl]cyclohexanecarboxylic acid and 4-[4-(3-chloro-4-hydroxybenzylamino)benzo[4,5]thieno[2,3-d]pyrimidin-2-yl]cyclohexanecarboxylic acid.

IT **313641-90-6P**
RL: IMF (Industrial manufacture); PNU (Preparation, unclassified); PREP (Preparation)
(preparation of benzo-annelated heterocycles by aromatization of tetrahydrobenzo-annelated heterocycles with hydrogenation catalyst in the presence of hydrogen acceptor)

RN 313641-90-6 CAPLUS

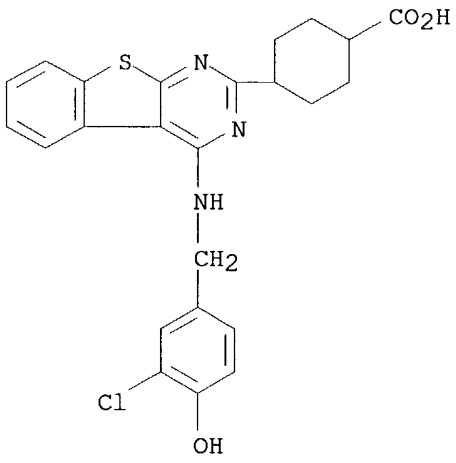
CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)



IT **313641-90-6DP**, alkyl esters
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of benzo-annelated heterocycles by aromatization of tetrahydrobenzo-annelated heterocycles with hydrogenation catalyst in the presence of hydrogen acceptor)

RN 313641-90-6 CAPLUS

CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:525954 CAPLUS
DOCUMENT NUMBER: 135:112007
TITLE: Pharmaceuticals containing 2-pyrrolidone as the dissolving solvent
INVENTOR(S): Schreder, Sven; Schamp, Karen; Wildner, Claudia
PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany
SOURCE: PCT Int. Appl., 34 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

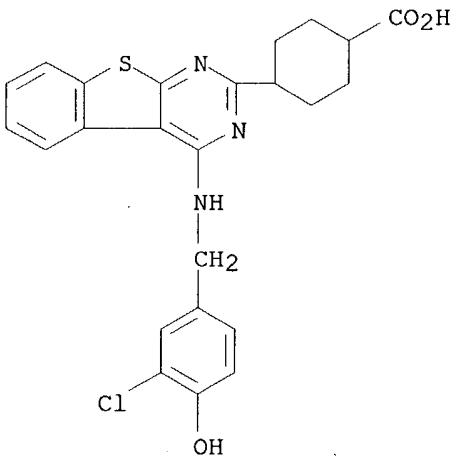
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001051089	A1	20010719	WO 2001-EP81	20010105
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1255565	A1	20021113	EP 2001-942307	20010105
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
US 2003153585	A1	20030814	US 2002-169396	20021101

PRIORITY APPLN. INFO.: DE 2000-10001020 A 20000113
WO 2001-EP81 W 20010105

AB The invention relates to pharmaceuticals containing at least 1 drug and 2-pyrrolidone as the dissolving solvent. Thus, a formulation was prepared from the ethanolamine salt of 4-[4-(3-chloro-4-methoxybenzylamino)benzo[4,5]thieno[2,3-d]pyrimidin-2-yl]cyclohexylcarboxylic acid ethanolamine salt 50, 2-pyrrolidone 100, and Gelucire 44/14 780 mg.

IT **313641-90-6**

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(pharmaceuticals containing pyrrolidone as dissolving solvent)
RN 313641-90-6 CAPLUS
CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:523501 CAPLUS
DOCUMENT NUMBER: 135:111982
TITLE: Pharmaceutical formulation containing (benzothienopyrimidinyl)cyclohexancarboxylate derivative
INVENTOR(S): Schreder, Sven; Wildner, Claudia; Schamp, Karen
PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany
SOURCE: Ger. Offen., 8 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10001021	A1	20010719	DE 2000-10001021	20000113
WO 2001051052	A1	20010719	WO 2001-EP82	20010105

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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

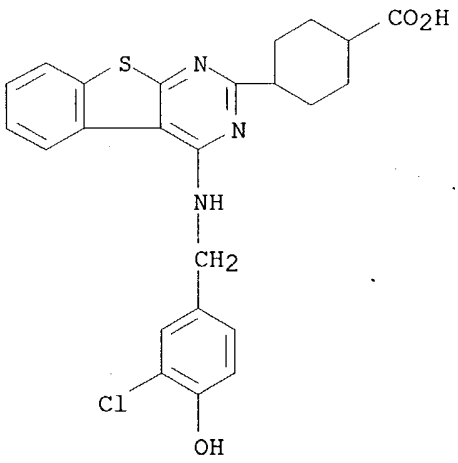
PRIORITY APPLN. INFO.: DE 2000-10001021 A 20000113
AB The invention concerns a pharmaceutical formulation containing 4-[4-(3-chloro-4-methoxybenzylamino)benzo[4,5]thieno[2,3-d]pyrimidin-2-

yl]cyclohexancarboxylic acid (I) or its salts and/or 4-[4-(3-chloro-4-hydroxybenzylamino)benzo[4,5]thieno[2,3-d]pyrimidin-2-yl]cyclohexancarboxylic acid or its salts, a substrate, a disintegrant and if necessary a glidant. Thus, a capsule formulation contained the ethanolamine salt of I 56.4, EtOH 900.0, microcryst. cellulose 143.3, highly-dispersed SiO2 7.2, and sodium carboxymethyl starch 5.0 mg.

IT **313641-90-6**
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(pharmaceutical formulations containing (benzothienopyrimidinyl)cyclohexancarboxylate)

RN 313641-90-6 CAPLUS

CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)



L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:	2000:897971 CAPLUS
DOCUMENT NUMBER:	134:56682
TITLE:	Preparation of benzothienopyrimidinealkanoates and analogs as cGMP phosphodiesterase inhibitors
INVENTOR(S):	Jonas, Rochus; Schelling, Pierre; Kluxen, Franz-Werner; Christadler, Maria
PATENT ASSIGNEE(S):	Merck Patent G.m.b.H., Germany
SOURCE:	Ger. Offen., 10 pp. CODEN: GWXXBX
DOCUMENT TYPE:	Patent
LANGUAGE:	German
FAMILY ACC. NUM. COUNT:	1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 19928146	A1	20001221	DE 1999-19928146	19990619
WO 2000078767	A1	20001228	WO 2000-EP5278	20000607

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1189907	A1	20020327	EP 2000-936871	20000607	
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO					
BR 2000011778	A	20020423	BR 2000-11778	20000607	
JP 2003502429	T2	20030121	JP 2001-504933	20000607	
AU 767273	B2	20031106	AU 2000-52204	20000607	
NO 2001006201	A	20011218	NO 2001-6201	20011218	
ZA 2002000440	A	20030417	ZA 2002-440	20020117	

PRIORITY APPLN. INFO.: DE 1999-19928146 A 19990619
WO 2000-EP5278 W 20000607

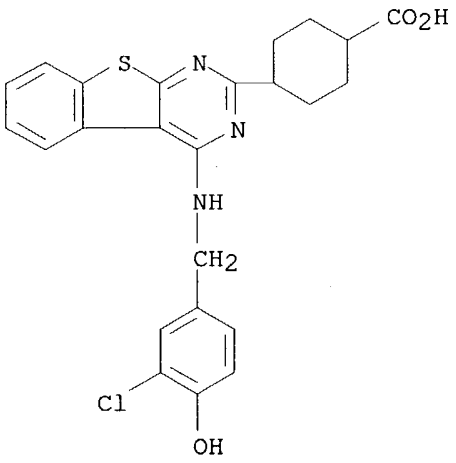
OTHER SOURCE(S): MARPAT 134:56682

AB R1CH2NHZR [Z = benzo[4,5]thieno[2,3-d]pyrimidine-4,2-diyl][I; R = Z1R7; R1 = (un)substituted Ph; R7 = CO2H, alkoxycarbonyl, cyano, (di)(alkyl)carbamoyl; Z1 = alk(en)ylene, cycloalkylene, phenylene, etc.] were prepared as cGMP phosphodiesterase inhibitors (no data). Thus, e.g., I (R = CH2CH2CO2H, R1 = C6H3(OMe)Cl-4,3) was prepared

IT **313641-90-6P 313641-91-7P 313641-92-8P**
313641-93-9P 313641-94-0P 313641-95-1P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of benzothienopyrimidinealkanoates and analogs as cGMP phosphodiesterase inhibitors)

RN 313641-90-6 CAPLUS

CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)

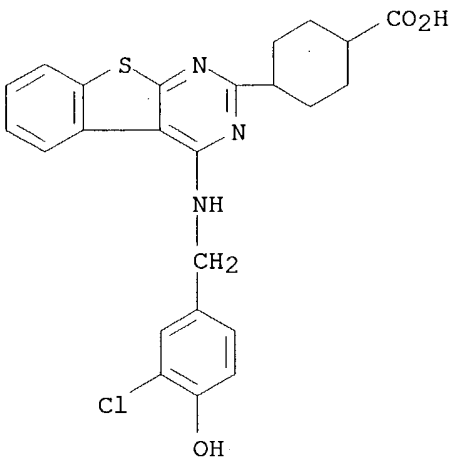


RN 313641-91-7 CAPLUS

CN Cyclohexanecarboxylic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]-, compd. with 2-aminoethanol (1:1) (9CI) (CA INDEX NAME)

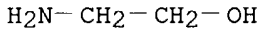
CM 1

CRN 313641-90-6
CMF C24 H22 Cl N3 O3 S

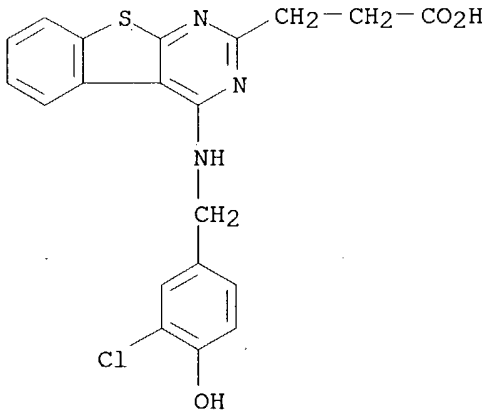


CM 2

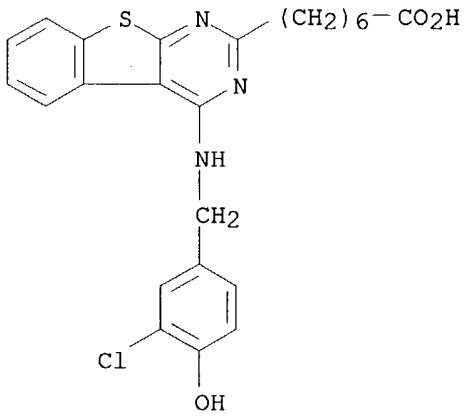
CRN 141-43-5
CMF C2 H7 N O



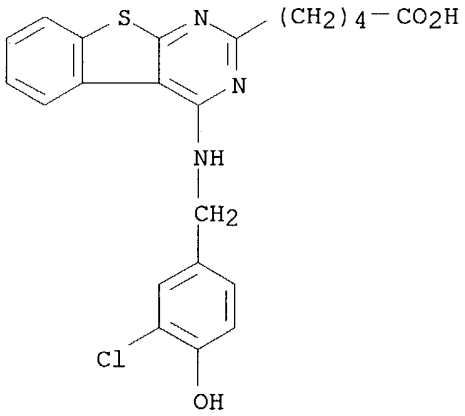
RN 313641-92-8 CAPLUS
CN [1]Benzothieno[2,3-d]pyrimidine-2-propanoic acid, 4-[[(3-chloro-4-hydroxyphenyl)methyl]amino]- (9CI) (CA INDEX NAME)



RN 313641-93-9 CAPLUS
CN [1]Benzothieno[2,3-d]pyrimidine-2-heptanoic acid, 4-[[(3-chloro-4-hydroxyphenyl)methyl]amino]- (9CI) (CA INDEX NAME)



RN 313641-94-0 CAPLUS
CN [1]Benzothieno[2,3-d]pyrimidine-2-pentanoic acid, 4-[[(3-chloro-4-hydroxyphenyl)methyl]amino]- (9CI) (CA INDEX NAME)



RN 313641-95-1 CAPLUS
CN Cyclohexaneacetic acid, 4-[4-[[(3-chloro-4-hydroxyphenyl)methyl]amino][1]benzothieno[2,3-d]pyrimidin-2-yl]- (9CI) (CA INDEX NAME)